

CLAIMS

1. A recombinant single chain polynucleotide comprising a region encoding the variable region of the light chain of an anti-G<sub>D2</sub> antibody linked to a region encoding the variable region of the heavy chain of an anti-G<sub>D2</sub> antibody.
2. The recombinant polynucleotide of claim 1, further comprising a region encoding an additional protein.
3. The recombinant polynucleotide of claim 1, wherein the additional protein is streptavidin.
4. The recombinant polynucleotide of claim 1, wherein the additional protein is a drug-converting enzyme.
5. A recombinant single chain peptide comprising the variable region of the light chain of an anti-G<sub>D2</sub> antibody linked to the variable region of the heavy chain of an anti-G<sub>D2</sub> antibody.
6. The peptide according to claim 5, wherein the peptide is labeled with a radiolabel.
7. The peptide according to claim 6, wherein the radiolabel is <sup>99m</sup>Tc.
8. The peptide according to claim 5, wherein the peptide further comprises a drug-converting enzyme.
9. The peptide according to claim 5, wherein the peptide further comprises streptavidin.

10. The peptide according to claim 5, wherein the peptide further comprises CD8.

11. T cells expressing a recombinant single chain peptide comprising the variable region of the light chain of an anti- $G_{D2}$  antibody linked to the variable region of the heavy chain of an anti- $G_{D2}$  antibody.

12. A method for assaying for the presence of cells expressing  $G_{D2}$  in tissue comprising combining the tissue with a recombinant single chain peptide comprising the variable region of the light chain of an anti- $G_{D2}$  antibody linked to the variable region of the heavy chain of an anti- $G_{D2}$  antibody and a detectable label.

13. A method for targeted delivery of a therapeutic agent to cells expressing  $G_{D2}$  in tissue comprising combining the tissue with a recombinant single chain peptide comprising the variable region of the light chain of an anti- $G_{D2}$  antibody linked to the variable region of the heavy chain of an anti- $G_{D2}$  antibody and a therapeutic or pre-therapeutic moiety.

14. The method according to claim 13, wherein the pre-therapeutic moiety is a pro-drug converting enzyme.

15. The method according to claim 13, wherein the pre-therapeutic moiety is streptavidin.

16. The method according to claim 13, wherein the therapeutic moiety is a toxin.